

**Amendments to the Claims:**

This listing of claims will replace all prior versions, and listing, of claims in the application:

**Listing of Claims:**

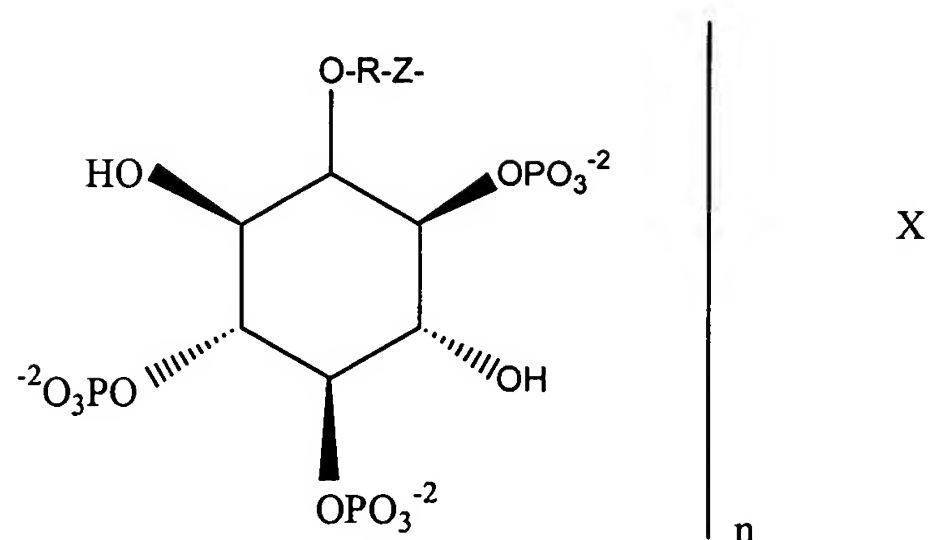
1. (Currently Amended): A protein binding assay for measuring inositol 1,4,5-triphosphate (IP<sub>3</sub>) in a sample employing as reagents a conjugate of IP<sub>3</sub> and a detectable label joined through a bond or linker at the 2-hydroxyl position of said IP<sub>3</sub>, and as a binding protein a truncated extracellular portion of an inositol 1,4,5-triphosphate receptor (IP<sub>3</sub>R) having at least about 200 times the affinity for IP<sub>3</sub> than the intact IP<sub>3</sub>R, wherein said conjugate and IP<sub>3</sub> in the sample compete for binding to said binding protein and the amount of bound or unbound conjugate will be related to the number of binding proteins bound by IP<sub>3</sub> in said sample, said method comprising:  
  
combining in an assay medium said sample, said conjugate and said binding protein and incubating said mixture for sufficient time for ~~any~~ complex formation of IP<sub>3</sub> and said conjugate ~~to bind to~~ with said binding protein; and  
  
detecting the bound or unbound label as a measure of the IP<sub>3</sub> present in the sample.
2. (Original): A protein binding assay according to Claim 1, wherein said assay is in a homogeneous format.
3. (Original): A protein binding assay according to Claim 1, wherein said sample is a cellular lysate, and wherein said cellular lysate has been treated to block kinases and phosphatases and prepare said sample for said assay.
4. (Original): A protein binding assay according to Claim 1, wherein said binding protein is of not more than about 600 amino acids and comprises at least amino acids 226 – 578 of the mouse IP<sub>3</sub>R Type 1.

5. (Original): A protein binding assay according to Claim 1, wherein said label is an enzyme fragment for enzyme complementation.
6. (Original): A protein binding assay according to Claim 1, wherein said binding protein is a fusion protein of up to about 1.5kD amino acids.
7. (Original): A protein binding assay according to Claim 1, wherein said label is a fluorescer.
8. (Original): A method according to Claim 1, wherein the order of addition of reagents is: (a) combining said sample with said binding protein; and (b) adding said conjugate, with incubating after (a) and (b).
9. (Withdrawn): A protein binding assay for measuring  $IP_3$  in a sample using a homogeneous format, employing as reagents a conjugate of  $IP_3$  and an ED of from 37 to 60 amino acids derived from  $\beta$ -galactosidase joined through a linker at the 2-hydroxyl position, and a truncated extracellular portion of an  $IP_3R$  having at least about 200 times the affinity for  $IP_3$  than the intact  $IP_3R$ , said method comprising:  
  
combining in an assay medium assay components in the following order: said sample, said binding protein, said conjugate and EA, and incubating after each combining for sufficient time for complex formation between said assay components;  
  
adding substrate for said  $\beta$ -galactosidase; and  
  
detecting the turnover of said  $\beta$ -galactosidase of said substrate as a measure of the  $IP_3$  present in the sample.
10. (Withdrawn): A protein binding assay for measuring  $IP_3$  in a sample using a homogeneous format, employing as reagents a conjugate of  $IP_3$  and a fluorescer joined through a linker at the 2-hydroxyl position, and a truncated extracellular portion of an  $IP_3R$  having at least about 200 times the affinity for  $IP_3$  than the intact  $IP_3R$ , said method comprising:

combining in an assay medium assay components: said sample, said binding protein, and said conjugate, and incubating for sufficient time for complex formation between said assay components; and

detecting the change in fluorescence polarization as a measure of the IP<sub>3</sub> present in the sample.

11. (Withdrawn): A method according to Claim 10, wherein said linker is an aliphatic group of from 4 to 20 carbon atoms.
12. (Currently amended): A method according to ~~Claim 9~~ Claim 10, wherein said fluorescer emits at a wavelength greater than about 500 nm.
13. (Withdrawn): A method according to Claim 10, wherein said fluorescer has a polarizability of less than about 60mP.
14. (Withdrawn): A protein binding assay for measuring IP<sub>3</sub> in a sample employing as reagents a conjugate of IP<sub>3</sub> and a detectable label joined through a bond or linker at the 2-hydroxyl position, and a truncated extracellular portion of an IP<sub>3</sub>R having at least about 200 times the affinity for IP<sub>3</sub> than the intact IP<sub>3</sub>R, said method comprising:  
  
combining in an assay medium said sample, said conjugate, said binding protein and a chemical reductant and incubating said mixture for sufficient time for any IP<sub>3</sub> and said conjugate to bind to said binding protein; and  
  
detecting the bound or unbound label as a measure of the IP<sub>3</sub> present in the sample.
15. (Withdrawn): A protein binding assay according to Claim 14, wherein said chemical reductant is a thiol.
16. (Withdrawn): A compound of the formula:



wherein:

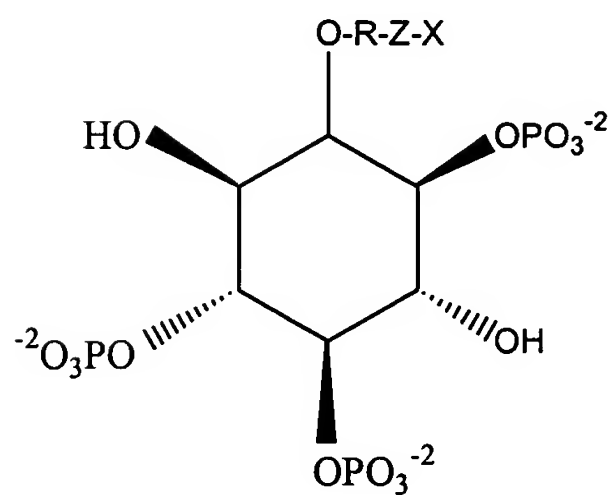
R is a neutral linking group of from 4 to 20 carbon atoms bonded to the oxygen through a saturated carbon atom or carbonyl;

Z is a functionality for linking X to the oxygen at the 2-position;

X is an enzyme donor fragment of  $\beta$ -galactosidase of from 27 to 60 amino acids;  
and

n is 1 or 2.

17. (Withdrawn): A compound of the formula:



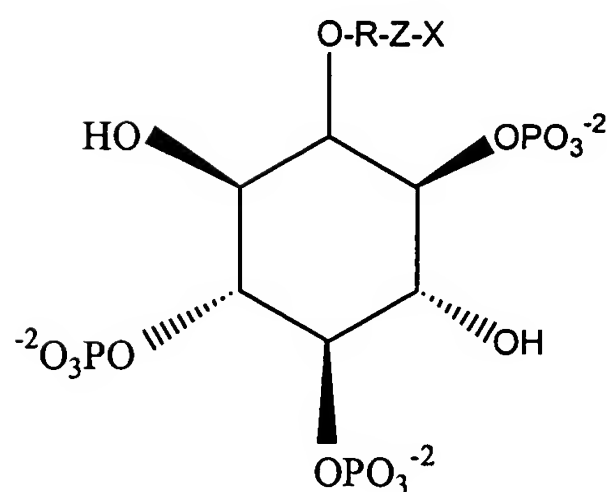
wherein:

R is a neutral linking group of from 2 to 20 carbon atoms bonded to the oxygen through a saturated carbon atom;

Z is a functionality for linking X to the oxygen at the 2-position; and

X is a fluorescer.

18. (Currently amended): A kit comprising a compound ~~according to Claim 17,~~  
~~enzyme acceptor for said enzyme donor and a truncated extracellular portion of an~~  
~~IP<sub>3</sub>R having at least about 200 times the affinity for IP<sub>3</sub> than the intact IP<sub>3</sub>R.~~



wherein:

R is a neutral linking group of from 2 to 20 carbon atoms bonded to the oxygen through a saturated carbon atom;

Z is a functionality for linking X to the oxygen at the 2-position; and

X is a fluorescer, enzyme acceptor for said enzyme donor and a truncated extracellular portion of an IP<sub>3</sub>R having at least about 200 times the affinity for IP<sub>3</sub> than the intact IP<sub>3</sub>R.

19. (Canceled).

20. (Withdrawn): A kit for performing an IP<sub>3</sub> assay comprising a conjugate of IP<sub>3</sub> and a detectable label joined through a bond or linker at the 2-hydroxyl position, a truncated extracellular portion of an IP<sub>3</sub>R having at least about 200 times the affinity for IP<sub>3</sub> than the intact IP<sub>3</sub>R and instructions for performing said assay.
21. (Withdrawn): A kit according to Claim 20, further comprising a thiol reductant.